Amend the claims as follows:

1\64.7 (Amended Once)\ A therapeutic method for controlling thrombosis and decreasing blood hypercoagulation and hemorrhaging risks in a patient which comprises administering to the patient in an antithrombotic effective amount, a composition which comprises a therapeutically acceptable carrier and heparinic mucopolysaccharide fractions having constituents of a molecular weight not in excesse of about 10,000 daltons, which fractions have (1) a mixture of lower molecular weight fractions in the range of about 2,000 to about 4,000 daltons with higher molecular weight fractions of a molecular weight in the range of about 4,000 to about 10,000 daltons, (2) a Yin-Wessler of at least 40, and (3) a ratio of Yin-Wessler to USP titer in the range of 3 to 5, and the physiologically acceptable salts thereof, which mixture of fractions have improved antithrombotic activity in vivo which is higher than that of heparin and a whole anticoagulation activity lower than that of heparin, and said method controlling thrombosis by selectively inhibiting coagulation factor Xa while also having a whole anticoagulation effect which is slower and lower than that of heparin.

A therapeutic composition for controlling thrombosis and decreasing hemorrhaging and of blood hypercoagulation risks which comprises a therapeutically

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having constituents of a molecular weight not in excess of about 10,000 daltons, which fractions have (1) a mixture of lower molecular weight fractions in the range of about 2,000 to about 4,000 daltons with higher molecular weight fractions of a molecular weight in the range of about 4,000 to about 10,000 daltons, (2) a Yin-Wessler of at least 40, and (3) a ratio of Yin-Wessler to USP titer in the range of 3 to 5, and the physiologically acceptable salts thereof, which mixture of fractions have improved antithrombotic activity in vivo which is higher than that of heparin and a whole anticoagulation activity lower and slower than that of heparin.

Add the following claims:

constituents of a molecular weight not in excess of about 10,000 daltons, which fractions have (1) a mixture of lower molecular weight fractions in the range of about 2,000 to about 4,000 daltons with higher molecular weight fractions of a molecular weight in the range of about 4,000 to about 10,000 daltons, (2) a Yin-Wessler of at least 40, and (3) a ratio of Yin-Wessler to USP titer in the range of 3 to 5, which mixture of fractions have improved antithrombotic activity in vivo which is higher than that of heparin and a whole anticoagulation activity lower than that of heparin, and the physiologically acceptable salts thereof.

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170. The heparinic mucopolysaccharides of claim 169 in which the lower molecular fractions have a USP titer less than about 10 units per mg.

wherein the molecular weight is not in excess of about 8,000 daltons.

The heparinic mucopolysaccharides of claim, 169 which have a USP titer of about 45 units per mg, a Yin-Wessler titer of about 160 units/mg and a ratio of Yin-Wessler to USP titer of about 3.55.

173. The heparinic mucopolysaccharides of claim, 169 in which fractions below 4,000 have a ratio of Yin-Wessler to USP titer which is at least 10.

The heparinic mucopolysaccharides of claim 169 in which fractions of about 4,000 have a ratio of Yin-Wessler to USP titer higher than 11, and the Yin-Wessler is at least 900 units per mg.

The heparinic mucopolysaccharides of claim 169 in which fractions have a Yin-Wessler to USP titer ratio higher than 60 and a Yin-Wessler of at least 1,300 units per mg.

wherein fractions above 4,000 have a USP titer not exceeding about 15 units per mg and a Yin-Wessler titer in the range of about 99 to about 160 units per mg.

wherein the fractions have a ratio of a Yin-Wessler to USP titer is in the range of about 13 to about 16.

wherein fractions have a USP titer that does not exceed about 6 units per mg, a Yin-Wessler titer not less than about 44 units per mg and the ratio of Yin-Wessler to USP titers if about at least 9.

179. The heparinic mucopolysaccharides of claim 169 having low molecular weight fractions with specific affinity for antithrombin III.

180. The heparinic mucopolysaccharides of claim, 169 in which fractions have 8 to 12 monosaccharide units corresponding to a molecular weight ranging from about 2,500 to 3,800.

wherein fractions have a molecular weight range of about 2,000 to about 8,000.

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192. The heparinic mucopolysaccharide fractions of claim 169, which are soluble in an aqueous-alcoholic medium, and insoluble in pure alcohol.

183. A therapeutic composition which presents less risks than heparin of blood hypercoagulation and of a host hemorrhaging, which composition has improved antithrombotic activity (anti- X_a activity) and improved selectivity with respect to anti- X_a activity than heparin in vivo and a lower and slower anticoagulation activity than heparin, and which composition comprises a therapeutically acceptable carrier and an antithrombotic effective amount of heparinic mucopolysaccharide fractions having constituents of a molecular weight not in excess of about 10,000 daltons, which fractions have (1) a mixture of lower molecular weight fractions in the range of about 2,000 to about 4,000 daltons with higher molecular weight fractions of a molecular weight in the range of about 4,000 to about 10,000 daltons, (2) a Yin-Wessler of at least 40, and (3) a ratio of Yin-Wessler to USP titer in the range of 3 to 5, which mixture of fractions have improved antithrombotic activity in vivo which is higher than that of heparin and a whole anticoagulation activity lower than that of heparin, and the physiologically acceptable salts thereof.

The therapeutic composition of claim 183 in which the lower molecular fractions of the heparinic mucopoly-saccharides have a USP titer less than about 10 units per mg.

185. The therapeutic composition of claim 183 in which the molecular weight of the heparinic mucopolysaccharides is not in excess of about 8,000 daltons.

186. The therapeutic composition of claim 183 in which the heparinic mucopolysaccharides have a USP titer of about 45 units per mg, a Yin-Wessler titer of about 160 units/mg and a ratio of Yin-Wessler to USP titer of about 3.55.

187. The therapeutic composition of claim 183 in which fractions of the heparinic mucopolysaccharides below 4,000 have a ratio of Yin-Wessler to USP titer which is at least 10.

188. The therapeutic composition of claim 183 in which heparinic mucopolysaccharides have fractions of about 4,000 which have a ratio of Yin-Wessler to USP titer higher than 11, and the Yin-Wessler is at least 900 units per mg.

the therapeutic composition of claim 183 in which fractions of the heparinic mucopolysaccharides have a Yin-Wessler to USP titer ratio higher than 60 and a Yin-Wessler of at least 1,300 units per mg.

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190. The therapeutic composition of claim 183 wherein fractions of the heparinic mucopolysaccharides above 4,000 have a USP titer not exceeding about 15 units per mg and a Yin-Wessler titer in the range of about 99 to about 160 units per mg.

Tel. The therapeutic composition of claim 190 wherein the fractions of the heparinic mucopolysaccharides have a ratio of a Yin-Wessler to USP titer is in the range of about 13 to about 16.

192. The therapeutic composition of claim 183 wherein fractions of the heparinic mucopolysaccharides have a USP titer that does not exceed about 6 units per mg, a Yin-Wessler titer not less than about 44 units per mg and the ratio of Yin-Wessler to USP titers if about at least 9.

193. The therapeutic composition of claim 183 wherein the heparinic mucopolysaccharides have low molecular weight fractions with specific affinity for antithrombin III.

194. The therapeutic composition of claim 187 wherein fractions of the heparinic mucopolysaccharides have 8 to 12 monosaccharide units corresponding to a molecular weight ranging from about 2,500 to 3,800.

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